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LOGINID: SSPTAEXB1618

PASSWORD:

TERMINAL (ENTER 1, 2, 3, OR ?):2

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NEWS
NEWS
     2 OCT 02
                 CA/CAplus enhanced with pre-1907 records from Chemisches
                 Zentralblatt
NEWS 3 OCT 19
                 BEILSTEIN updated with new compounds
NEWS 4 NOV 15
                 Derwent Indian patent publication number format enhanced
NEWS 5
         NOV 19
                 WPIX enhanced with XML display format
NEWS 6
         NOV 30 ICSD reloaded with enhancements
NEWS 7 DEC 04 LINPADOCDB now available on STN NEWS 8 DEC 14 BEILSTEIN pricing structure to change
NEWS 9 DEC 17 USPATOLD added to additional database clusters
NEWS 10 DEC 17 IMSDRUGCONF removed from database clusters and STN
NEWS 11 DEC 17 DGENE now includes more than 10 million sequences
NEWS 12 DEC 17 TOXCENTER enhanced with 2008 MeSH vocabulary in
                 MEDLINE segment
NEWS 13 DEC 17 MEDLINE and LMEDLINE updated with 2008 MeSH vocabulary
NEWS 14 DEC 17 CA/CAplus enhanced with new custom IPC display formats
NEWS 15 DEC 17
                 STN Viewer enhanced with full-text patent content
                 from USPATOLD
NEWS 16 JAN 02
                 STN pricing information for 2008 now available
NEWS 17 JAN 16
                 CAS patent coverage enhanced to include exemplified
                 prophetic substances
NEWS 18 JAN 28 USPATFULL, USPAT2, and USPATOLD enhanced with new
                 custom IPC display formats
NEWS 19 JAN 28 MARPAT searching enhanced
NEWS 20 JAN 28 USGENE now provides USPTO sequence data within 3 days
                 of publication
NEWS 21 JAN 28 TOXCENTER enhanced with reloaded MEDLINE segment
NEWS 22 JAN 28 MEDLINE and LMEDLINE reloaded with enhancements
NEWS 23 FEB 08 STN Express, Version 8.3, now available
NEWS 24 FEB 20 PCI now available as a replacement to DPCI
NEWS 25 FEB 25 IFIREF reloaded with enhancements
NEWS 26 FEB 25
                 IMSPRODUCT reloaded with enhancements
NEWS 27 FEB 29
                 WPINDEX/WPIDS/WPIX enhanced with ECLA and current
                 U.S. National Patent Classification
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NEWS EXPRESS FEBRUARY 08 CURRENT WINDOWS VERSION IS V8.3, AND CURRENT DISCOVER FILE IS DATED 20 FEBRUARY 2008

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NEWS LOGIN Welcome Banner and News Items
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=> file caplus
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FILE COVERS 1907 - 11 Mar 2008 VOL 148 ISS 11 FILE LAST UPDATED: 10 Mar 2008 (20080310/ED)

Effective October 17, 2005, revised CAS Information Use Policies apply. They are available for your review at:

http://www.cas.org/infopolicy.html

=> s bis(4-hydroxyaryl)alkane MISSING OPERATOR 'BIS(4-HYDROXYA' The search profile that was entered contains terms or nested terms that are not separated by a logical operator.

=> s 4-hydroxyaryl 5812114 4 1737 HYDROXYARYL

2 HYDROXYARYLS

1738 HYDROXYARYL

(HYDROXYARYL OR HYDROXYARYLS)

L1 107 4-HYDROXYARYL

(4(W)HYDROXYARYL)

=> s adduct

86064 ADDUCT 68783 ADDUCTS

L2 124304 ADDUCT

(ADDUCT OR ADDUCTS)

=> s 11 and 12

```
T.3
             9 L1 AND L2
=> s phenol
        259014 PHENOL
        125481 PHENOLS
L4
        324247 PHENOL
                 (PHENOL OR PHENOLS)
=> s 13 and 14
             8 L3 AND L4
=> d bib abs hitstr 1-8
L_5
     ANSWER 1 OF 8 CAPLUS COPYRIGHT 2008 ACS on STN
ΑN
     2007:592329 CAPLUS
     147:10341
DN
    Methods for increasing the mean particle size of 2-hydrocarbyl-3,3-
ΤI
     bis(hydroxyaryl)phthalimidines
     Ganesan, Balakrishnan; Nadkarni, Pradeep Jeevaji
ΙN
     General Electric Company, USA
PA
     U.S. Pat. Appl. Publ., 15 pp.
SO
     CODEN: USXXCO
DT
     Patent
LA
    English
FAN.CNT 1
                       KIND
                                DATE
                                           APPLICATION NO.
     PATENT NO.
                                                                   DATE
                        ----
                        A1
PΙ
     US 2007123712
                                20070531
                                            US 2005-288912
                                                                    20051129
                        В2
     US 7329720
                                20080212
                        A2
     WO 2007064623
                                20070607
                                            WO 2006-US45506
                                                                    20061128
                               20070726
     WO 2007064623
                         А3
            AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH,
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             KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK,
             MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO,
             RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT,
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             IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ,
             CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH,
             GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
             KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA
                              20051129
PRAI US 2005-288912
                      A
    MARPAT 147:10341
OS
     A method for increasing a mean particle size of a 2-hydrocarby1-3,3-
AB
     bis(hydroxyaryl)phthalimidine is provided. The method comprises forming a
     mixture comprising a feedstream of the 2-hydrocarbyl-3,3-bis(4-
     hydroxyaryl)phthalimidine, and a solvent composition comprising an organic
     solvent and water, wherein the organic solvent is capable of at least
     partially dissolving the 2-hydrocarbyl-3,3-bis(hydroxyaryl)phthalimidine
     and forming an adduct with the 2-hydrocarbyl-3,3-
bis(hydroxyaryl)phthalimidine. Then the mixture is heated at a temperature and
     for a time effective to decompose the adduct and form a
     2-hydrocarbyl-3,3-bis(hydroxyaryl)phthalimidine product having a mean
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producing polymers.

RE.CNT 19 THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

bis(hydroxyaryl)phthalimidines with increased particle size are useful for

particle size greater than 5 μ . The 2-hydrocarbyl-3,3-

```
2004:740283 CAPLUS
AN
DN
     141:245239
     Process for recovering an adduct of a bis(4-
ΤI
     hydroxyaryl) alkane and a phenolic compound
     Patrascu, Emil; Frey, Johann-Wilhelm; Hagel, Manfred
IN
PA
     Dow Global Technologies, Inc., USA; Dow Deutschland Inc.
SO
     PCT Int. Appl., 18 pp.
     CODEN: PIXXD2
     Patent
DT
     English
LA
FAN.CNT 1
     PATENT NO.
                       KIND DATE
                                         APPLICATION NO.
                                                                 DATE
                        ____
                                           _____
                                20040910 WO 2004-US1118
PΙ
     WO 2004076394
                        A1
                                                                  20040116
         W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH,
             CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD,
             GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,
             LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI
         RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE,
             BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU,
            MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN,
             GQ, GW, ML, MR, NE, SN, TD, TG
     EP 1597224
                                           EP 2004-702992
                         A1
                               20051123
                                                                   20040116
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK
                            20060329
                       Α
                                         CN 2004-80004859
                                                             20040116
     CN 1753856
                         Τ
                              20060810
     JP 2006518377
                                            JP 2006-502852
                                                                   20040116
     US 2006224025
                        A1 20061005
                                           US 2005-541779
                                                                   20050711
     IN 2005CN01964
                        A 20070727
                                            IN 2005-CN1964
                                                                   20050818
                        Р
PRAI US 2003-448918P
                             20030222
                               20030221
                        W
     WO 2004-US1118
     A process for recovering a solid adduct of a bis(4-
AΒ
     hydroxyaryl)alkane and a phenolic compound from a suspension
     comprising the addict, comprises the steps of: (a) supplying the
     suspension to a rotary filter; (b) filtering the supplied suspension in
     the rotary filter to retain adduct as an adduct cake;
     (c) pre-drying the adduct cake with an inert gas; (d) washing
     the pre-dried adduct cake; (e) optionally drying the washed
     adduct cake; and (f) discharging the washed adduct cake
     from the rotary filter. Thus, a pure bis(4-hydroxyaryl
     )alkane is obtained through the adduct recovered when it is
    melted and the phenolic compound is distilled off.
RE.CNT 2
             THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
             ALL CITATIONS AVAILABLE IN THE RE FORMAT
     ANSWER 3 OF 8 CAPLUS COPYRIGHT 2008 ACS on STN
L5
     2004:398398 CAPLUS
ΑN
     141:156901
DN
     Oxidative nucleophilic substitution of hydrogen in nitrobenzenes with
ΤI
     2-phenylpropionic esters
     Makosza, Mieczyslaw; Surowiec, Marek; Paszewski, Maciej
ΑU
CS
     Institute of Organic Chemistry, Polish Academy of Sciences, Warsaw, PL-01
     224, Pol.
     ARKIVOC (Gainesville, FL, United States) (2004), (2), 172-180
SO
     CODEN: AGFUAR
     URL: http://www.arkat-usa.org/ark/journal/2004/Zwanenburg/BZ-
     975E/975E.pdfhttp://www.arkat-usa.org/ark/journal/2004/Zwanenburg/BZ-
     975E/975E.pdf
    Arkat USA Inc.
PΒ
    Journal; (online computer file)
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ANSWER 2 OF 8 CAPLUS COPYRIGHT 2008 ACS on STN

L5

AB Several alkyl 2-phenyl-2-(4-nitroaryl)propionates, e.g. I, and 2-phenyl-2-(4-hydroxyaryl)propionates, e.g. II, were prepared, in 66% and 73% yield, by oxidation of σ H adducts with KMnO4 and dimethyldioxirane, which were generated in situ from alkyl 2-phenylpropionates, e.g iso-Pr 2-phenylpropanoate and nitroarenes, e.g. 3-bromonitrobenzene.

RE.CNT 21 THERE ARE 21 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 4 OF 8 CAPLUS COPYRIGHT 2008 ACS on STN

AN 2001:449831 CAPLUS

DN 135:46601

TI Separation of bis(4-hydroxyaryl)alkanes and aromatic hydroxy compounds from bis(4-hydroxyaryl)alkane/hydroxyarene adducts in a desorber.

IN Neumann, Rainer; Heydenreich, Frieder; Prein, Michael; Lanze, Rolf; Boediger, Michael

PA Bayer A.-G., Germany

SO Ger. Offen., 6 pp. CODEN: GWXXBX

DT Patent

LA German

FAN.CNT 1

| PI DE 19961566 A1 20010621 DE 1999-19961566 19991 WO 2001046104 A1 20010628 WO 2000-EP12324 20001 W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, | | | | |
|---|----------|--|--|--|
| W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, | 19991220 | | | |
| | 20001207 | | | |
| CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, | CN, | | | |
| | HR, | | | |
| HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, | LT, | | | |
| LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, | RU, | | | |
| SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, | VN, | | | |
| YU, ZA, ZW | | | | |
| RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, | CY, | | | |
| DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, | BF, | | | |
| BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG | | | | |
| BR 2000016494 A 20020917 BR 2000-16494 20001 | 20001207 | | | |
| EP 1242349 A1 20020925 EP 2000-991585 20001 | 20001207 | | | |
| R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, | PT, | | | |
| IE, SI, LT, LV, FI, RO, MK, CY, AL, TR | | | | |
| JP 2003518048 T 20030603 JP 2001-546618 20001 | 20001207 | | | |

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TW 526190 B 20030401 TW 2000-89127151 20001219
IN 2002MN00705 A 20040228 IN 2002-MN705 20020530
US 2002183562 A1 20021205 US 2002-149906 20020617
                           B2 20050719
A 20030128 MX 2002-PA6090
      US 6919487
     MX 2002PA06090
                                                                            20020619
RR 786460 B1 20071217

KR 2007110447 A 20071116

PRAI DE 1999-19961566 A 19991220

WO 2000-EP12324 W 20001207

KR 2002-707867 A3 20020619
                                                KR 2002-707867
KR 2007-723912
                                                                             20020619
                                                                             20071018
      Use of a desorber optionally in series with a distillation unit for separation
AΒ
of
      bis(4-hydroxyaryl)alkanes [specifically
      2,2-bis(4-hydroxyphenyl)propane, BPA] and aromatic hydroxy compds. from bis(
      4-hydroxyaryl)alkane/arylhydroxy adducts is
      claimed. Desorption is carried out in a desorber consisting of
      tube-bundle heat exchangers; interstices between the heat exchanger pipes
      are filled with ceramic balls (steatite). An inert gas (N2 or O2) is fed
      through the desorber at a flow of 100-300~\mathrm{m3} per \mathrm{m3}~\mathrm{BPA/PhOH}
      adducts at 160-230^{\circ}. BPA is recovered as bottom product in
      the desorber and collected in a withdrawal tank. Separation of BPA from
      BPA/PhOH adducts gave BPA in a purity of >99.5% with PhOH
      content of <50 ppm.
L5
      ANSWER 5 OF 8 CAPLUS COPYRIGHT 2008 ACS on STN
      2001:449826 CAPLUS
AN
     135:46600
DN
      separation and purification of bis(4-hydroxyaryl
ΤI
     )alkanes using a vacuum drum filter
     Neumann, Rainer; Lanze, Rolf; Heydenreich, Friedrich; Boediger, Michael;
ΤN
     Prein, Michael
     Bayer A.-G., Germany
PA
SO
     Ger. Offen., 6 pp.
      CODEN: GWXXBX
DT
     Patent
LA
     German
FAN.CNT 1
                      KIND DATE APPLICATION NO. DATE
     DE 19961521
WO 2001046105
                            A1 20010621 DE 1999-19961521
A1 20010628 WO 2000-EP12323
PΙ
          W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
               CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR,
               HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT,
               LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU,
               SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN,
               YU, ZA, ZW
          RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
               DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
               BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
      BR 2000016505
                             Α
                                    20020827 BR 2000-16505
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      EP 1242350
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Т3
      JP 2003518049
                                    20030603
                                                JP 2001-546619
                                                                             20001207
     ES 2218277 T3 20041116 ES 2000-990667
TW 568901 B 20040101 TW 2000-89127150
IN 2002MN00733 A 20040313 IN 2002-MN733
MX 2002PA06089 A 20030128 MX 2002-PA6089
US 2003038094 A1 20030227 US 2002-149905
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                                                                          20001207
20001219
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                                                                            20020619
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US 6906227 B2 20050614
HK 1054920 A1 20060106 HK 2003-107259 20031009
PRAI DE 1999-19961521 A 19991220 WO 2000-EP12323 W 20001207
      Adducts of bis(4-hydroxyaryl)alkanes (prepared
AB
      by acid-catalyzed reaction of aromatic hydroxy compds. with ketones) with
      hydroxyarenes are separated and purified by continuous filtration in a
      rotating vacuum drum filter. The drum filter contains filter cells
      including a suction zone, a washing zone, a dry suction zone, an aeration
      zone, and optionally a filter cake withdrawal zone and a cloth filter
      washing zone. The crystals (filter cake) are separated in an amount of 800
kq/h
      and washed in the washing zone with 50-150% PhOH (filter cake basis) at
      45-70°. Process conditions (e.g. drum speed, filter cake
      thickness, circulation N2) are set so that the residual moisture content
      of the filter cake is <30%. Purified adduct crystals are melted
      on a heating spiral and collected in collecting tanks. Purification of
      2,2-bis(4-hydroxyphenyl)propane (BPA) according to the process gave BPA
      crystals in a purity of >99% and with PhOH content of <50 ppm.
L5
      ANSWER 6 OF 8 CAPLUS COPYRIGHT 2008 ACS on STN
ΑN
      2000:254116 CAPLUS
DN
      132:280883
ΤI
      Manufacture of bis(4-hydroxyaryl)alkanes
      Kuehling, Steffen; Lanze, Rolf; Neumann, Rainer; Heydenreich, Frieder; Van
ΙN
      Osselaer, Tony
      Bayer A.-G., Germany
PΑ
SO
      Ger. Offen., 4 pp.
      CODEN: GWXXBX
DT
      Patent
LA
     German
FAN.CNT 1
      PATENT NO. KIND DATE APPLICATION NO. DATE
      DE 19848026 A1 20000420 DE 1998-19848026 19981017
TW 517046 B 20030111 TW 1999-88116896 19991001
WO 2000023410 A1 20000427 WO 1999-EP7358 19991005
PΙ
           W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU,
                CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL,
                IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD,
                MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK,
                SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW
           RW: GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE,
                DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF,
                CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
      AU 9960893 A 20000508 AU 1999-60893
BR 9914607 A 20010703 BR 1999-14607
                                                                                19991005
                                                                                19991005
      EP 1121339
                              A1
                                                   EP 1999-947458
                                     20010808
                                                                                19991005
                              B1 20030129
      EP 1121339
           R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
                IE, SI, LT, LV, FI, RO
MD 2001000161 A 20010930
MD 2705 B2 20050228
JP 2002527497 T 20020827
ES 2190253 T3 20030716
MX 2001PA03769 A 20010731
US 6384288 B1 20020507
US 2002055661 A1 20020509
PRAI DE 1998-19848026 A 19981017
WO 1999-EP7358 W 19991005
US 2001-807645 A3 20010416
                                                    MD 2001-20010161
                                                                                 19991005

      JP 2000-577138
      19991005

      ES 1999-947458
      19991005

      MX 2001-PA3769
      20010411

      US 2001-807645
      20010416

      US 2002-37995
      20020103
```

Bis(4-hydroxyaryl)alkanes are separated from their AΒ adducts with aromatic OH compds. by (a) passing an inert gas through molten adducts and stripping the phenols at 150-230°, (b) removing the stripped phenols from the inert gas by condensation, and (c) purifying, compressing and recirculating the inert gas into the step (a). Thus, bisphenol A with Hazen color number 8 was obtained by use of N for removing PhOH from a molten 60/40% bisphenol A/PhOH mixture as described above.

RE.CNT 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

- L_5 ANSWER 7 OF 8 CAPLUS COPYRIGHT 2008 ACS on STN
- ΑN 2000:115788 CAPLUS
- DN 132:166709
- ΤI Recovery of bis(4-hydroxyaryl)alkanes with increased purity from their adducts with phenols
- Kuehling, Steffen; Lanze, Rolf; Neumann, Rainer; Heydenreich, Frieder; Van ΤN Osselaer, Tony; Fennhoff, Gerhard

- PΑ Bayer A.-G., Germany
- Ger., 4 pp. SO
 - CODEN: GWXXAW
- DTPatent
- LA German
- FAN.CNT 1

| L MIV • | | PATENT NO. | | | | | KIND | | DATE | | | PLI | CAT | | DATE | | | | |
|---------|----------|---|-------------------|-------------------|-------------------|-------------------|-------------------|---|-------------------|----------------------------------|----------------|----------------------------|-------------------|-------------------|-------------------|-------------------|------------|------------|------------|
| PI | | 19840110 2000014044 | | | | | | | | | | | | | | | | | |
| | | ₩: | CZ, IN, MG, | DE, IS, MK, | DK, JP, MN, | DM, KE, MW, | EE, KG, MX, | AZ, ES, KP, NO, UA, | FI, KR, NZ, | GB, KZ, PL, | GI LC PI |), C, I, | GE, LK, RO, | GH, LR, RU, | GM, LS, SD, | HR, LT, SE, | HU, LU, | ID, LV, | IL, MD, |
| | | R₩: | GH, ES, | GM, FI, | KE, FR, | LS, GB, | MW, GR, | SD, IE, | SL, IT, | SZ, LU, | UG MC | ġ, Ċ, | ZW, NL, | AT, PT, | BE, | CH, | , | , | , |
| | | - , - , - , | | | , | A | , | N, ML, MR, NE, SN, TD, TG 20000327 AU 1999-58540 | | | | | | | | | | | |
| | EP | | | | A1 | | | | | | | | | | | | | | |
| | EΡ | | AT, | BE, | CH, | DE, | DK, | ES, | | | GF | ٦, | IT, | LI, | LU, | NL, | SE, | MC, | PT, |
| | JP ES | IE, SI, LT, 2002524434 2179674 | | | T | | 2002 2003 | | | JP 2000-568804 ES 1999-946008 | | | | | | | | | |
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AΒ Bis(4-hydroxyaryl)alkanes with increased purity and reduced purity variation are manufactured from their adducts with aromatic hydroxy compds. which are prepared by acid-catalyzed conversion of the aromatic hydroxy compds. with ketones. The crystalline adducts are treated with aerosol dispersions of aqueous alkali metal hydroxide solns. with variable concentration (0.005-0.015%), and then separated from phenols by distillation Thus, treating continuously crystalline bisphenol A/PhOH adduct with aerosol dispersion of aqueous NaOH solution via gas phase while monitoring (GC) the amount of impurities (isopropenylphenol, isopropenylphenol dimer and trisphenol) and increasing the NaOH concentration in the aerosol when the

total impurity concentration exceeded 100 ppm, gave bisphenol A of higher

and reduced the purity variation.

THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD RE.CNT 7 ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 8 OF 8 CAPLUS COPYRIGHT 2008 ACS on STN

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2,2-Bis(4-hydroxyaryl)propanes

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SO 5 pp.

DT Patent

Unavailable LA

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DE 116100: KIND DATE DATE APPLICATION NO. 19640116 DE 1961-U7974 DE 1161284 19610428 PΙ GB 974982 GB

19600506 PRAI US

Title compds. were prepared by the reaction of 1 mole of allene, propyne, or mixts. thereof with 3-20 moles of an appropriate phenol having a sterically unhindered, reactive para H atom, at 30-125° (preferably $55-60^{\circ}$ in the presence of an insol., strongly acid cation exchange resin containing 0.01-0.5, especially 0.175, acid equivalent/mole of phenol, such as a sulfonated styrene-divinylbenzene copolymer or a phenol -formaldehyde sulfonic acid resin), under nearly water-free conditions. Thus, a stirred mixture of 564 g. molten phenol and 250 g. (0.875)acid equivalent) Dowex 50~W cation exchange resin, dried to a water-content of <2%, was heated to 50° , 40 q. of a 70:30% mixture of propyne:allene added over 3.5 hrs. through a gas-inlet tube placed below the surface of the liquid, the mixture filtered, the filter cake washed with 250 cc. molten phenol, the filtrate and the filter cake washed with 250 cc. molten phenol, and the filtrate and washings combined and distilled at 1 mm. to a final residue temperature of 200° to yield 183 g. crude 2,2-bis(4-hydroxyphenyl)propane (I) in the residue. The crude product was purified by heating it with >1:1 ratio of phenol:crude product at $37-95^{\circ}$. The by-products and a small amount of I are soluble, while most of I forms a crystalline 1:1 adduct with phenol. The adduct was filtered off or centrifuged, washed with phenol , then heated to remove the phenol, which was recycled to the reaction vessel, as were the filtrate and washings containing by-products, and unreacted olefins. The residue consisted of very pure I. Because an equilibrium between I and by-products of the reaction occurred and remained constant under constant reaction conditions, no accumulation of by-products took place, and the process displayed an efficiency of >99%. Other examples showed effect of reaction variables on yield, however, the above example detailed reflected optimum conditions.